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IC. AMENDMENTS TO THE CLAIMS

Please enter the amendments to claims 1 and 12-14, as shown below.

- (Currently amended) A method for diagnosing Alzheimer's Disease (AD) [[AD]] in a
 living individual, the method comprising detecting a level of carboxyl-terminal truncated apolipoprotein
 E (apoE) in an aqueous biological sample from the individual, wherein a level of carboxyl-terminal
 truncated apoE that is significantly higher than the level present in a normal control indicates that the
 individual has AD.
 - 2. (Original) The method of claim 1, wherein the biological sample is blood.
 - 3. (Original) The method of claim 1, wherein the biological sample is serum.
- (Original) The method of claim 1, wherein the carboxyl-terminal truncated apoE has a molecular weight of about 14-20 kDa.
- (Original) The method of claim 1, wherein the carboxyl-terminal truncated apoE comprises amino acids 244-260 of apoE.
 - 6. (Original) The method of claim 1, wherein apoE is apoE4.
 - (Original) The method of claim 1, wherein apoE is apoE3.
 - (Original) The method of claim 1, wherein apoE is a mixture of apoE3 and apoE4.
 - (Canceled)
- 10. (Previously presented) The method of claim 1, further comprising detecting a level of full length apolipoprotein E (apoE) in the biological sample from the individual; wherein a ratio of the level of carboxyl-terminal truncated apoE compared to the level of full length apoE in the biological sample

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that is greater than a ratio associated with a control biological sample from an individual not having Alzheimer's Disease is indicative of a diagnosis of Alzheimer's Disease.

- (Previously presented) The method of claim 10, wherein the carboxyl-terminal truncated apoE has a molecular weight of about 14-20 kDa.
- 12. (Currently amended) The method of claim 10, wherein a ratio of the level of carboxylterminal truncated apoE compared to the level of full length apoE in the biological sample that is at least
 25% greater than a ratio associated with a control biological sample from an individual not having
 Alzheimer's Disease is indicative of a diagnosis of Alzheimer's Disease the ratio is greater than about
 1.5.
- 13. (Currently amended) The method of claim 10, wherein a ratio of the level of carboxyl-terminal truncated apoE compared to the level of full length apoE in the biological sample that is at least 50% greater than a ratio associated with a control biological sample from an individual not having Alzheimer's Disease is indicative of a diagnosis of Alzheimer's Disease the ratio is greater than about 2.
- 14. (Currently amended) The method of claim 10, wherein a ratio of the level of carboxyl-terminal truncated apoE compared to the level of full length apoE in the biological sample that is at least 2-fold greater than a ratio associated with a control biological sample from an individual not having Alzheimer's Disease is indicative of a diagnosis of Alzheimer's Disease the ratio is greater than about 3.
- 15. (withdrawn) A kit for diagnosing Alzheimer's Disease, the kit comprising an antibody that binds to carboxyl-terminal truncated apolipoprotein E (apoE) and instructions for using the antibody for diagnosing Alzheimer's Disease.
- (withdrawn) The kit of claim 15, further wherein the antibody is attached to a solid support.
- (withdrawn) The kit of claim 15, further comprising an antibody that specifically binds to a carboxyl-terminal portion of full-length apoE.

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(withdrawn) The kit of claim 15, wherein the instructions for the diagnosis of
 Alzheimer's Disease direct the use of the kit to detect carboxyl-terminal truncated apoE in serum.

- 19. (Previously presented) The method of claim 1, wherein the biological sample is plasma.
- (Previously presented) The method of claim 1, wherein the biological sample is cerebrospinal fluid.
- (withdrawn) The kit of claim 17, wherein the antibody that specifically binds to a carboxyl-terminal portion of apoE binds specifically to an epitope within amino acids 270-299 of apoE.
 - 22. (withdrawn) The kit of claim 16, wherein the solid support is a test strip.